

Supplementary Figure 7. Read length distribution profiles vary between sRNA-seq datasets. We

downloaded publicly available sRNA-seq datasets from the NCBI SRA database at <a href="https://www.ncbi.nlm.nih.gov/sra">https://www.ncbi.nlm.nih.gov/sra</a> (accessions summarized in Supplementary Table 8) and analyzed the read length profiles of reads between 15 and 40 bases in length. The x-axis shows number of reads and the y-axis the read length in bases. We examined datasets from the following samples: (A) *H. vulgare* infected with *B. hordei* at 0, 24, and 48 hpi (Hunt et al. 2019); (B) *H. vulgare* under salt stress (Deng et al. 2015) and aluminium stress (Wu et al. 2018), respectively; (C) Triticum aestivum (wheat) after infection with *B. graminis* f.sp. tritici at 12 hpi and under 40 °C heat stress (Xin et al. 2011); (D) *T. aestivum* infected with *Zymoseptoria tritici* at 12 dpi (Ma et al. 2019); (E) *T. aestivum* under 37 °C heat stress, continuous light stress, or UV treatment stress (Ragupathy et al. 2016); (F) *Glycine max* (soybean) during nodulation with the bacterial species *Bradyrhizobium japonicum* at 10 and 20 days after inoculation (Ren et al. 2019); (G) *Arabidopsis thaliana* and *Phaseolus vulgaris* (common bean) during infection with *Sclerotinia sclerotiorum* (Derbyshire et al. 2019); (H) *A. thaliana* after infection with *Verticillium dahliae* and the *V. dahliae* mutant aly1 aly2 (Zhu et al. 2022); (I) *A. thaliana* infected with *Hyaloperonospora arabidopsidis* at 3, 4, and 7 dpi (Dunker et al. 2020); (J) Botrytis cinerea cultivated in vitro (Weiberg et al. 2013); (K) *A. thaliana* infected with *B. cinerea* at 24, 48, and 72 hpi (Weiberg et al. 2013).